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1995:532272 CAPLUS
AN
    122:258639
DN
     Ketoglutaric acid for controlling microorganisms and seaweeds in the
ΤI
     culture medium of laver
    Okuzono, Kazuhiko
IN
     Daiichi Seimo Kk, Japan
PA
     Jpn. Kokai Tokkyo Koho, 4 pp.
SO
     CODEN: JKXXAF
     Patent
DT
     Japanese
LA
     ICM A01N037-42
ICS A01G033-02
IC
     5-2 (Agrochemical Bioregulators)
FAN.CNT 1
                                          APPLICATION NO. DATE
                     KIND DATE
     PATENT NO.
     ______
                                          _____
                   A2 19950228
                                         JP 1993-200843
                                                          19930812
     JP 07053310
PΙ
     Ketoglutaric acid at \geq 0.1 % by wt./vol. is effective in
AΒ
     controlling microorganisms harmful to laver, and controlling
     undesirable seaweeds interfering with the growth of laver.
     Ketoglutaric acid may be used in conjunction with nitrogen source
     like ammonium nitrate, and with acids like citric acid, and HCl.
     laver culture microbicide carboxylic acid
ΙT
     Algicides
     Bactericides, Disinfectants, and Antiseptics
     Laver
        (control of microorganisms and seaweeds in culture medium of
        laver with ketoglutaric acid and)
IT
        (control of microorganisms and seaweeds in culture medium of
        laver with ketoglutaric acid and acids)
     50-21-5, Lactic acid, biological studies 57-13-6, Urea, biological
IT
     studies 64-18-6, Formic acid, biological studies 64-19-7, Acetic
     acid, biological studies 76-03-9, Trichloroacetic acid, biological
     studies 77-92-9, Citric acid, biological studies
     Monochloroacetic acid, biological studies 79-43-6, Dichloroacetic
     acid, biological studies 87-69-4, Tartaric acid, biological
     studies 110-15-6, Succinic acid, biological studies 110-16-7,
     Maleic acid, biological studies 110-17-8, Fumaric acid, biological
     studies 526-95-4, Gluconic acid 6484-52-2, Ammonium nitrate,
     biological studies 6915-15-7, Malic acid 7558-80-7, Monosodium
     phosphate 7631-99-4, Sodium nitrate, biological studies
     7647-01-0, Hydrochloric acid, biological studies 7664-38-2,
     Phosphoric acid, biological studies 7664-93-9, Sulfuric acid,
     biological studies 7697-37-2, Nitric acid, biological studies
     7722-76-1, Mono-ammonium phosphate 7778-77-0, Monopotassium
     phosphate 12125-02-9, Ammonium chloride, biological
             15421-51-9, Inositol phosphate
     RL: AGR (Agricultural use); BAC (Biological activity or effector,
     except adverse); BIGL (Biological study); USES (Uses)
        (control of microorganisms and seaweeds in culture medium of
        laver with ketog... aric acid and)
ΙT
     328-50-7
     RL: AGR (Agricultural use); BAC (Biological activity or effector,
     except adverse); BIOL (Biological study); USES (Uses)
        (for controlling microorganisms and seaweeds in culture medium of
        laver:
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MEDLINE

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83074524
ΑN
                PubMed ID: 6816216
     83074524
DN
     Change in subunit composition of the iron protein of nitrogenase
ΤI
     from Rhodospirillum rubrum during activation and inactivation of
     iron protein.
     Preston G G; Ludden P W
AII
     BIOCHEMICAL JOURNAL, (1982 Sep 1) 205 (3) 489-94.
SO
     Journal code: 9YO; 2984726R. ISSN: 0264-6021.
     ENGLAND: United Kingdom
CY
     Journal; Article; (JOURNAL ARTICLE)
DT
     English
LΑ
     Priority Journals
FS
     198301
EM
     Entered STN: 19900317
FD
     Last Updated on STN: 19970203
     Entered Medline: 19830119
     The subunit composition of the Fe protein of nitrogenase from
AB
     Rhodospirillum rubrum during activation and inactivation was
     investigated. It was found that the upper subunit (on gel
     electrophoresis) of the two-subunit Fe protein was converted into
     the lower subunit during activation in vitro. When the Fe protein
     was inactivated in vivo by the addition of NH4Cl and
     alpha-oxoglutarate to the cells, a phosphate-labelled upper band
     appeared. During activation in vitro by the activating enzyme, some
     of the phosphate of the upper band remained with the protein and
     appeared in the lower band. Activations in vitro were performed on
     inactive Fe protein obtained from cells grown with glutamate as the
     nitrogen source. Both native and oxygen-denatured Fe protein
     exhibited the loss of upper band during treatment with activating
     enzyme.
     Check Tags: Support, Non-U.S. Gov't; Support, U.S. Gov't, Non-P.H.S.
CT
      Ammonium Chloride: PD, pharmacology
     *Bacterial Proteins: ME, metabolism
      Chromatography, Gel
      Enzyme Activation: DE, drug effects
      Ketoglutaric Acids: PD, pharmacology
      *Metalloproteins: ME, metabolism
      *Nitrogenase: ME, metabolism
      Nonheme Iron Proteins
      *Rhodospirillum rubrum: EN, enzymology
     12125-02-9 (Ammonium Chloride); 328-50-7 (alpha-ketoglutaric acid)
RN
      0 (Bacterial Proteins); 0 (Ketoglutaric Acids); 0 (Metalloproteins);
CN
      0 (Nonheme Iron Proteins); EC 1.18.6.1 (Nitrogenase)
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WPI

- Agent contg. keto-glutaric acid - used for controlling and eliminating algae ΤI and bacteria from cultured layer AB

J07053310 Agent contains ketoglutaric acid.

- Also claimed is an agent contg. at least 1 of ammonium nitrate, ammonium chloride, monoammonium phosphate, urea, sodium nitrate and nitric acid as a nitrogen source, monopotassium phosphate, monosodium phosphate and inositol 6-phosphate (phytic acid) as a phosphorus source, in addn. to at least 0.1 wt./vol.% ketoglutaric acid.

Also claimed is an agent contg. citric acid, malic acid, tartaric acid, succinic acid, gluconic acid, acetic acid, mono-, di- or tri-chloroacetic acid, lactic acid, maleic acid, fumaric acid, formic acid, hydrochloric acid, phosphoric acid, nitric acid and sulphuric acid, in addn. to at least 0.1 wt. /vol.% ketoglutaric acid.

- USE/ADVANTAGE - The agent is useful for eliminating algae e.g. Enteromorpha adhered on a laver culturing net, and for eliminating bacteria belonging to genus Phythium or Olpidiopsis. The agent has strong bactericidal activity to bacteria belonging to genus Pythium or Olpidiopsis. The time required for eliminating the Enteromorpha is short.

- In an example, a 0.5 w/v sea water soln. of ketoglutaric acid was prepd. and Enteromorpha was immersed in the soln. for 5 mins., then taken out from the soln., washed with sea water and static cultured was carried out after 2 days the Enteromorpha was discoloured or decoloured.(Dwg.0/0)

- JP7053310 A 19950228 DW199517 A01N37/42 004pp PN

- JP19930200843 19930812 PR

- (DAII-N) DAIICHI SEIMO KK PA

- C05-A01A C05-A01B C05-B01P C05-B02A2 C05-B02A3 C05-C C10-A07 C10-A13C C10-MC C02 C14-A01 C14-A05 D05-H01 D09-A01C

DC - C03 D16 P13

IC - A01G33/02 ;A01N37/42

AN - 1995-128197 [17]

PAJ

- EXTERMINATING AGENT AGAINST MISCELLANEOUS ALGAE AND DISEASE INJURIES OF ΤI

- PURPOSE: To provide a medicine capable of exterminating miscellaneous algae AΒ and a bacterium or a fungus of the genus Pythium, Olpidiopsis, etc., causing red rot or chytrid blight in an ultrashort time as compared with that of other organic acids.

- CONSTITUTION: This exterminating agent against miscellaneous algae and disease injuries of cultured laver contains >=0.1wt./vol.% ketoglutaric acid as the active ingredient. At least one selected from the group consisting of ammonium nitrate, ammonium chloride, monoammonium phosphate, urea, sodium nitrate and nitric acid as the nitrogen source and monopotassium phosphate, monosodium phosphate, phosphoric acid and inositol 6-phosphate as the phosphorus source can be contained in the exterminating agent. Furthermore, at least one selected from the group consisting of citric acid, malic acid, tartaric acid, succinic acid, gluconic acid, acetic acid, chloroacetic acid, lactic acid, maleic acid, fumaric acid, formic acid, hydrochloric acid, phosphoric acid, nitric acid and sulfuric acid can be used in combination so as to enhance exterminating effects. The fungus of the genus Pythium causing red rot can be exterminated in 1min. and green laver can be exterminated in 5min. by immersing the laver in a solution of seawater containing 0.5wt./vol. % ketoglutaric acid.

PN - JP7053310 A 19950228

PD- 1995-02-28 ABD - 19950630

ABV - 199505

ΑP - JP19930200843 19930812

PA - DAIICHI SEIMO KK IN OKUZONO KAZUHIKO

- A01N37/42 ;A01G33/02